Claims:

- 1. Pharmaceutical preparation that contains at least one emulsifier, at least one auxiliary emulsifier and/or solvent as well as at least one lipid, characterized in that the mass ratio of emulsifier to auxiliary emulsifier and/or solvent (Smix) is 1:1 to 9:1 and the total lipid proportion is > 0% (m/m), whereby this preparation at least partially inhibits at least one intestinal enzyme and/or at least one intestinal efflux system.
- Pharmaceutical preparation according to claim 1, wherein the Smix is
 3:1 to 9:1.
- 3. Pharmaceutical preparation according to claim 3, wherein the Smix is9:1.
- 4. Pharmaceutical preparation according to at least one of claims 1-3, wherein the total lipid proportion is 10-50% (m/v).
- 5. Pharmaceutical preparation according to at least one of claims 1-4, whereby intestinal enzymes originate from the group of 17β-hydroxy-steroid-dehydrogenase or the cytochrome monooxygenases and intestinal efflux systems from the group of P-glycoproteins.
- 6. Pharmaceutical preparation according to at least one of claims 1 to 5, wherein the emulsifier contains PEG-40-hydrogenated castor oil (Cremophor®RH40), PEG-35 castor oil (Cremophor®EL) or PEG-400-monoricinoleate (Estax®54).
- 7. Pharmaceutical preparation according to at least one of claims 1-6, wherein the auxiliary emulsifier and/or the solvent contains glyceryl

- monocaprylate > 80% (m/m) (Imwitor®308) or diethylene glycol monoethyl ether (Transcutol®P).
- 8. Pharmaceutical preparation according to at least one of claims 1-7, wherein the lipid contains triglycerides, fatty oils or waxes.
- 9. Pharmaceutical preparation according to claim 8, wherein the triglyceride contains mid-chain triglycerides (Miglyol®).
- 10. Pharmaceutical preparation according to claim 8, wherein the fatty oil contains castor oil, olive oil, corn oil, soybean oil, sunflower oil, peanut oil, walnut oil or diestel oil.
- 11. Pharmaceutical preparation according to claim 8, wherein the wax contains ethyl oleate or isopropyl myristate.
- 12. Pharmaceutical preparation according to at least one of claims 1 to 11, wherein the preparation contains in addition at least one pharmaceutical substance.
- 13. Pharmaceutical preparation according to claim 12, wherein the pharmaceutical substance is lipophilic and/or water-insoluble or hydrophilic.
- 14. Pharmaceutical preparation according to claim 12 or 13, wherein at least one pharmaceutical substance is a substrate of at least one intestinal enzyme and/or an intestinal efflux system.
- 15. Pharmaceutical preparation according to claim 14, wherein at least one intestinal enzyme originates from the group of 17β-hydroxy-steroid-dehydrogenases and/or cytochrome-monooxygenases.

- 16. Pharmaceutical preparation according to claim 15, wherein at least one intestinal enzyme is 17β-HSD 2 and/or originates from the group of cytochrome P 450 3A-monooxygenases.
- 17. Pharmaceutical preparation according to claim 14, wherein at least one intestinal efflux system originates from the group of P-gp- transporter systems.
- 18. Pharmaceutical preparation according to one of claims 12 to 17, wherein at least one pharmaceutical substance is a steroid.
- 19. Pharmaceutical preparation according to claim 18, wherein the steroid in 17-position of the sterane skeleton contains a secondary, betaposition hydroxyl group.
- 20. Pharmaceutical preparation according to claim 18 or 19, wherein the steroid is an estrogen, an antiestrogen or an androgen.
- 21. Pharmaceutical preparation according to at least one of claims 18 to 20, wherein the steroid 11-α-hydroxynandrolone, 16-α-fluoroestradiol, 16-α-iodoestradiol, 16-β-fluoroestradiol, 2,4-dibromoestradiol, 2-chloroestradiol, 2-ethoxyestradiol, 2-fluoroestradiol, 2-hydroxyestriol, 2-methoxyestradiol, 2-methoxyestriol, 2-methoxyestradiol, 3-methoxyestriol, 4-bromoestradiol, 4-chloroestradiol, 4-fluoro-17β-estradiol, 4-hydroxyestradiol, 4-hydroxytestosterone, 4-methoxyestradiol, 5-β-androstan-17β-ol-3-one, 6-α-hydroxyestradiol, 3α, 17β-androstanediol, 3β,17β-androstanediol, androstanolone, androstenediol, bolanediol, bolazine, boldenone, clostebol, dacuronium

bromide, 17-deacetylpancuronium, dideactetylvecuronium, vecuronium, 17β-dihydroequilin, 5α-dihydro-19-nortestosterone, 16αbromo-7α-(N-butyl, N-methyl-undecanamide)-estra-1,3,5(10)-triene-3.17β-diol, 16α-chloro-7α-(N-butyl, N-methyl-undecanamide)-estra-1.3.5(10)-triene-3.17 β -diol, 16 α -iodo-7 α -(N-butyl, N-methylundecanamide)-estra-1,3,5(10)-triene-3,17β-diol, 16α-bromo-7α-(Nbutyl, N-methyl-undecanamide)-estra-1,3,5(10)-triene-3,17β-diol, epiestriol, epitiostanol, estetrol, estradiol, estradiol-3-glucuronide, estradiol-3-methylether, estradiol-3-sulfate, estradiol-3-benzoate, estradiol-3-hexahydrobenzoate, estramustine, estriol, estriol-3glucuronide, estriol-3-sulfate, estriol-16-glucuronide, estrynamine, 17β-hydroxy-6-methylene-androsta-1,4-dien-3-one, fulvestrant, 1hydroxy-17β-estradiol, 2-hydroxy-17β-estradiol, 4-hydroxy-17βestradiol, 6-hydroxy-17β-estradiol, 7-hydroxy-17β-estradiol, 15hydroxy-17β-estradiol, 18-hydroxy-17β-estradiol, 7-(N-butylundecanamide)-3,17 β -estra-1,3,5(10)-triene-3,17 β -diol, 7 α -(N-butylundecanamide)-3,17β-estra-1,3,5(10)-triene-3,17β-diol, estra-1,3,5(10)-triene-7 β -(N-butyl)undecanamide-3,17 β -diol, 7 α -(N-butyl, N-methyl-undecanamide)-estra-1,3,5(10)-triene-3,17 β -diol, inocoterone, estra-3-sulfamate-1,3,5(10),7-tetraene-3,17β-diol, cycloprop[14S,15β]-3',15-dihydro-estra-1,3,5(10)-triene-3,17β-diol, estra-1,3,5(10)-triene-3-sulfamate-17β-ol, mesterolone, methenolone,

16-methyleneestradiol, metogest, nandrolone, nisterime, norclostebol, 3-octyloxy-5α-androst-3-en-17β-ol, estradiol-17-phenylpropionateestradiol-benzoate mixture, 7-ethyl-nandrolone, 11β-chloromethylestra-3.17 β -diol, piperidinium-1- $\{(2\beta, 3\alpha, 5\alpha, 16\beta, 17\beta)-3, 17$ -dihydroxy-2-(1-piperidinyl)androstan-16-yl]-1-methyl-bromide, 17deacetylrocuronium, oxendolone, 11α-methoxy-7α-methyl-estra-3-17β-diol, quinestradol, 17β-hydroxy-7α-methyl-androst-5-en-3-one, 11 α -ethenyl-estra-3, 17 β -diol, 11 β -[4(dimethylamino)phenyl]-estra-3, 17β-diol, 7α -{4-[2-(dimethylamino)ethoxy]phenyl}-estra-3,17β-diol, 11B-{4-[(methylsulfonyl)oxy]phenyl}-estra-3,17B-diol, 11B-{4-[[5-[(4,4,5,5,5-pentafluoropentyl)sulfonyl]pentyl]oxy]phenyl}-estra-3,17βdiol, 17β-dihydroxy-9α-fluoro-11β-androsta-1,4-dien-3-one, stenbolone, cycloprop[14R,15\alpha]estra-3',15-dihydro-3-methoxy-1,3,5(10)-trien-17β-ol, cycloprop[14S,15β]estra-3',15-dihydro-3methoxy-1,3,5(10)-trien-17β-ol, testosterone, trestolone, trilostane, 13β-ethyl-8α-gona-1,3,5(10)-triene-3,16α,17β-triol, 13β-ethyl-8βgona-1,3,5(10)-triene-3,16 α ,17 β -triol, estra-2- $\{\text{tricyclo}[3.3.1.13,7]\text{decyl}\}$ -1,3,5(10)-triene-3,17 β -diol, ent-estradiol, 8 β -vinyl-estradiol, 11 β -fluoro-7 α -{5-[N-methyl-N-3-(4,4,5,5,5pentafluoropentylthio)-propylamino|pentyl}-estra-1,3,5(10)-triene-3,17 β -diol, 11 β -fluoro-7 α -{5-[methyl-(7,7,8,8,9,9,10,10,10nonafluorodecyl)amino]pentyl}estra-1,3,5(10)-triene-3,17β-diol, 11β-

fluoro- 17α -methyl- 7α - $\{5$ -[methyl-(8,8,9,9,9-pentafluorononyl)amino]pentyl} estra-1,3,5(10)-triene-3,17 β -diol, 17 β -hydroxy-14 α ,15 α methylene-androst-4-en-3-one, 17β -hydroxy- 7α -methyl- 14α , 15α methylene-androst-4-en-3-one, 4-chloro-17β-hydroxy-14α,15αmethylene-androst-4-en-3-one, 4,17β-dihydroxy-14α,15α-methyleneandrost-4-en-3-one, 17β-hydroxy-14α,15α-methylene-androsta-1,4dien-3-one, 4-chloro-17β-hydroxy-14α,15α-methylene-androsta-1,4dien-3-one, 4-chloro-17β-hydroxy-14α,15α-methylene-estr-4-en-3one, 7β-hydroxy-7α-methyl-14α,15α-methylene-estr-4-en-3-one, 17βhydroxy-14α,15α-methylene-estr-4-en-3-one, 4,17β-dihydroxy- $14\alpha,15\alpha$ -methylene-estr-4-en-3-one, 17β -hydroxy- $14\alpha,15\alpha$ methylene-estra-4,9,11-trien-3-one, 3-ethyl-17 β -hydroxy-14 α ,15 α methylene-gon-4-en-3-one, 17a-β-hydroxy-17a-homoandrosta-4,15dien-3-one, 1"-mesyl-17α-(trifluoromethyl)-1'Hpyrazol[4",5':2,3]androst-4-en-17 β -ol.

- Use of a pharmaceutical preparation according to at least one of claims

 1 to 21 for the production of a peroral pharmaceutical agent for
 inhibiting at least one intestinal enzyme and/or at least one intestinal
 efflux system.
- 23. Use of a pharmaceutical preparation according to claim 22, wherein at least one intestinal enzyme originates from the group of 17β-hydroxy-steroid-dehydrogenases and/or cytochrome-P450-monooxygenases.

- Use of a pharmaceutical preparation according to claim 23, wherein at least one intestinal enzyme is 17β -HSD 2 and/or originates from the group of cytochrome-P450-3A-monooxygenases.
- 25. Use of a pharmaceutical preparation according to claim 22, wherein at least one intestinal efflux system is a P-gp- transporter.
- Use of a pharmaceutical preparation according to at least one of claims22 to 25, wherein the pharmaceutical agent comes from the group oftherapeutic agents, prophylactic agents or diagnostic agents.
- 27. Process for increasing the bioavailability of pharmaceutical substances that are to be administered perorally, wherein a pharmaceutical preparation contains a pharmaceutical substance and is administered perorally according to at least one of claims 1 to 21.